

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

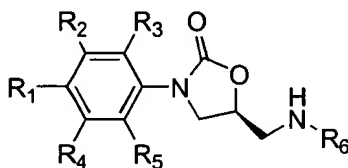
Claims 1-43 (Previously Cancelled).

Claims 44-48 (Currently Cancelled).

Claims 49-100 (Previously Cancelled).

Claim 101. (Currently Amended) A method of preparing combinatorial libraries of compounds of the formula Ib, comprising the steps of:

- a) attaching a plurality of aryl oxazolidinones to a plurality of solid supports;
 - b) functionalizing the 4-position of the aryl groups of the attached oxazolidinones to produce an R₁ substituent; and, optionally,
 - c) removing the oxazolidinones from the solid supports;
- wherein compounds of formula Ib have the structure:



wherein R₂, R₃, R₄ and R₅ are, independently, hydrogen alkyl, heteroalkyl, heteroaryl or an electron withdrawing group;

R₆ is acyl or sulfonyl; and

R₁ is one of the following functional groups:

C(O)NR₇R₈, wherein R₇ and R₈ are, independently, hydrogen, alkyl, heteroalkyl, aryl or heteroaryl;

C(O)OR₉, wherein R₉ is hydrogen, alkyl, heteroalkyl, aryl or heteroaryl;

C(O)R₁₀, wherein R₁₀ is hydrogen, alkyl, heteroalkyl, aryl or heteroaryl;

SR₁₁, wherein R₁₁ is hydrogen, alkyl, heteroalkyl, aryl or heteroaryl;

S(O)₂R₁₁, wherein R₁₁ is hydrogen, alkyl, heteroalkyl, aryl or heteroaryl;

S(O)R₁₁, wherein R₁₁ is hydrogen, alkyl, heteroalkyl, aryl or heteroaryl;

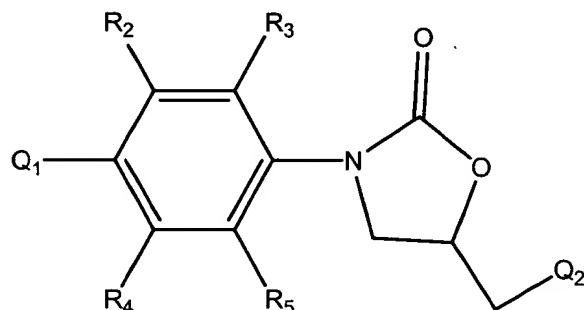
NR₁₂R₁₃, wherein R₁₂ and R₁₃ are, independently, hydrogen, acyl, sulfonyl, alkyl, heteroalkyl, aryl or heteroaryl;

2-oxazolyl, wherein R₁₄ is at the 4-position and R₁₅ is at the 5-position of the oxazolyl, and wherein R₁₄ and R₁₅ are, independently, hydrogen, alkyl, heteroalkyl, aryl, heteroaryl or an electron withdrawing group;

2-aminothiazolyl, wherein R₁₆ is at the 4-position and R₁₇ is at the 5-position of the thiazole, and wherein R₁₆ and R₁₇, are, independently, hydrogen, alkyl, heteroalkyl, aryl, heteroaryl or an electron withdrawing group;

CH₂NR₁₈R₁₉, wherein R₁₈ and R₁₉ are, independently, hydrogen, alkyl, heteroalkyl, aryl, heteroaryl, acyl or sulfonyl; and

wherein the aryl oxazolidinones in step a) comprise the structure



wherein R₂, R₃, R₄ and R₅ are defined above;

Q₁ is selected from -C(O)O-PG, -S-PG, -CH(OC₁₋₄alkyl)₂, where PG is a protecting group; and

Q₂ is N₃, wherein the resin includes a carbonyl containing moiety and attaching the plurality of aryl oxazolidinones to the plurality of solid supports in step a comprises converting Q₂ to an iminophosphorane or an amine and reacting the iminophosphorane or the amine with the carbonyl moiety of the resin to form an imine.

Cancel claims 102-107.

Claim 108. (Previously Added) The method of claim 101, wherein Q₁ is -C(O)O-PG.

Claim 109. (Previously Added) The method of claim 108, wherein functionalizing the 4-position of the aryl groups comprises converting the $-C(O)O-PG$ group into a $-C(O)NR_7R_8$, $-C(O)OR_9$, $-C(O)R_{10}$, $-NR_{12}R_{13}$, 2-oxazolyl, or 2-aminothiazolyl group.

Claim 110. (Previously Added) The method of claim 101, wherein Q_1 is $-S-PG$.

Claim 111. (Previously Added) The method of claim 110, wherein functionalizing the 4-position of the aryl groups comprises converting the $-S-PG$ group to a $-SR_{11}$, $-S(O)R_{11}$, or $-S(O)_2R_{11}$ group.

Claim 112. (Previously Added) The method of claim 101, wherein Q_1 is $-CH(OC_{1-4}alkyl)_2$.

Claim 113. (Currently Amended) The method of claim 112, wherein functionalizing the 4-position of the aryl groups comprises converting the $-CH(OC_{1-4}alkyl)_2$ group to a $[-CH_2NR_{12}R_{13}]$ $-CH_2NR_{18}R_{19}$ group.